



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁷ : C08F 10/06, 4/64</p>	<p>A1</p>	<p>(11) International Publication Number: WO 00/50476 (43) International Publication Date: 31 August 2000 (31.08.00)</p>
<p>(21) International Application Number: PCT/EP00/01736 (22) International Filing Date: 25 February 2000 (25.02.00) (30) Priority Data: 99103803.5 26 February 1999 (26.02.99) EP (71) Applicant (for all designated States except US): FINA RESEARCH S.A. [BE/BE]; Zone Industrielle C, B-7181 Seneffe (BE). (72) Inventor; and (75) Inventor/Applicant (for US only): DEMAIN, Axel [BE/BE]; Rue de Sart, 21, B-1457 Tourinnes-Saint-Lambert (BE). (74) Common Representative: FINA RESEARCH S.A.; Patent Dept., Zone Industrielle C, B-7181 Seneffe (BE).</p>		<p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
<p>(54) Title: USE AND PRODUCTION OF POLYPROPYLENE (57) Abstract Use of isotactic polypropylene homopolymers or copolymers in processes in which the polypropylene solidifies from a melt, wherein for enhanced speed of solidification of the polypropylene the polypropylene has a melt temperature and a crystallisation temperature not more than 50° less than the melt temperature resulting from the polypropylene having been produced using a metallocene catalyst component having the general formula (I): R" (C₉R₁R₂R₃) (C₉R'_a)_nMQ₂ wherein C₉ is a substituted cyclopentadienyl ring; C₉' is a substituted or unsubstituted fluorenyl ring; R" is a structural bridge imparting stereoregidity to the component; R₁ is a substituent on the cyclopentadienyl ring which is distal to the bridge, R₂ is a substituent on the cyclopentadienyl ring which is proximal to the bridge and positioned <i>non-vicinal</i> to the distal substituent, R₃ is a substituent on the cyclopentadienyl ring which is proximal to the bridge, each R'_a is the same or different and is hydrocarbyl having 1 to 20 carbon atoms in which oSn≤8; M is a Group IVB transition metal or vanadium and each Q is hydrocarbyl having 1 to 20 carbon atoms or is a halogen.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

USE AND PRODUCTION OF POLYPROPYLENE

The present invention relates to the use of homopolymers and copolymers of propylene prepared with a metallocene catalyst component in applications which require a low melting temperature, a high crystallisation temperature and a high transparency. The present invention also relates to a process for producing isotactic polypropylene homopolymers.

It is known in the art to obtain polypropylene with low melting temperature by inserting comonomers in the polymer chain during polymerisation. With Ziegler-Natta catalysts, the addition of ethylene (or other) comonomers in the growing chains of polypropylene during polymerisation gives rise to a random propylene copolymer that is characterised by a lower melting point, a lower flexural modulus, lower rigidity and higher transparency than the homopolymers of propylene. The comonomers generate defects in the polymer chain which impede the growth of thick crystalline structures and reduce the degree of crystallinity of the overall polymer. The comonomers are not evenly distributed in the polymer chains. Among the many comonomers that can be used in the copolymerisation process, ethylene and butene have been most frequently utilised. It has been observed that the melting temperature of the propylene copolymers is reduced by about 6°C per wt% of inserted ethylene in the copolymer chain or by about 3°C per wt% of inserted butene.

The addition of comonomer in industrial polymerisation processes has however other impacts than just decreasing the melting temperature of the polypropylene; it has both economical and technical impacts.

These known random propylene copolymers also suffer from the technical problem that the crystallisation temperature is

relatively low. This is technically disadvantageous when the polypropylene is being processed since the low crystallisation temperature increases the cycle time of any process where the polypropylene is being solidified from the melt. With the lower crystallisation temperatures, the period for solidification is longer, thereby increasing the cycle time for injection moulding, injection blow moulding and extrusion blow moulding and decreasing the production line speed in film, tube, profile or pipe extrusion.

EP-A-0881236 in the name of Fina Research S.A. and EP-A-0537130 in the name of Fina Technology, Inc. each disclose a metallocene catalyst component for use in producing isotactic polypropylene.

EP-A-0870779 in the name of Fina Technology, Inc. discloses metallocene catalysts for producing a blend of iso- and syndiotactic polypropylene.

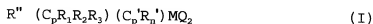
EP-A-0742227 in the name of Fina Technology, Inc. discloses a metallocene compound for producing hemisotactic polypropylene.

EP-A-0905173 discloses the production of biaxially oriented metallocene-based polypropylene films.

Despite the disclosures of these prior Fina patent specifications, there is a need in the art for polypropylene having not only relatively low melting temperature but also relatively high crystallisation temperature which enables the polypropylene to be used more readily in processes requiring the polypropylene to be solidified from the melt using shorter cycle times or higher film speeds.

Accordingly, the present invention provides use of isotactic polypropylene homopolymers or copolymers in processes in which the polypropylene solidifies from a melt, wherein for enhanced speed of solidification of the polypropylene the polypropylene has a melt temperature and a crystallisation temperature not

more than 50°C less than the melt temperature resulting from the polypropylene having been produced using a metallocene catalyst component having the general formula:



wherein C_p is a substituted cyclopentadienyl ring; C_p' is a substituted or unsubstituted fluorenyl ring; R'' is a structural bridge imparting stereorigidity to the component; R_1 is a substituent on the cyclopentadienyl ring which is distal to the bridge, which distal substituent comprises a bulky group of the formula XR^* , in which X is chosen from Group IVA, and when $a=3$ each R^* is the same or different and chosen from hydrogen or hydrocarbyl of from 1 to 20 carbon atoms, or when $a=2$ one R^* is chosen from hydrogen or hydrocarbyl of from 1 to 20 carbon atoms and the other different R^* is chosen from a substituted or unsubstituted cycloalkyl where X is a carbon atom in the cycloalkyl ring, R_2 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and positioned *non-vicinal* to the distal substituent and is hydrogen or of the formula $YR\#$, in which Y is chosen from Group IVA, and each $R\#$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, R_3 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and is a hydrogen atom or is of the formula $ZR\$$, in which Z is chosen from Group IVA, and each $R\$$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, each R_n' is the same or different and is hydrocarbyl having 1 to 20 carbon atoms in which $0 \leq n \leq 8$; M is a Group IVB transition metal or vanadium and each Q is hydrocarbyl having 1 to 20 carbon atoms or is a halogen.

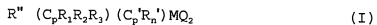
The metallocene catalyst component may be employed either alone or in a mixture of one or more metallocene catalyst components.

In accordance with the invention, it has now been found that homopolymers and copolymers of propylene obtained using these metallocene catalysts have characteristics which are similar to and better than those exhibited by random polypropylene (PP) copolymers produced using Ziegler-Natta catalysts. These desired characteristics are not only a low melting temperature and a high transparency, but also a high crystallisation temperature. The polymers usable in accordance with the invention can also tend to have higher rigidity as compared to random PP copolymers produced using Ziegler-Natta catalysts and higher flexibility than homopolymers produced using Ziegler-Natta catalysts. As well as having high transparency, the preferred polymers for use in the invention also have low haze.

The present invention is predicated on the discovery by the inventor that the use of particular metallocene catalysts enables polypropylene homopolymers, or polypropylene copolymers with a small amount of comonomer, to have a combination of relatively low melting temperature and relatively high crystallisation temperature which reduces the cycle time for processing the polymer from the melt, for example in injection moulding and injection or extrusion blow moulding. The amount of comonomer in the copolymers is not more than 25wt%, typically less than 10wt%, more preferably less than 5wt% and yet more preferably less than 3wt%. Typical comonomers are ethylene and butene, but other alpha-olefins may be employed. For extrusion processes, such as pipe, tube or profile extrusion and for film production, the higher crystallisation temperature permits higher line speeds. The film may be produced by casting, a tenter frame process or blowing. Other polyolefin processing methods exist for which the present invention has utility in which the polypropylene is processed from the melt. The combination of a low melting temperature and a high crystallisation temperature provides a reduced temperature "window" between those two temperatures, which enables the polypropylene to be more easily and more

quickly processed when the polypropylene is processed in the melt and then solidified from the melt.

Thus the present invention further provides a process for producing an isotactic homopolymer of propylene having a melt temperature of from 139 to 144°C and a difference between the melt temperature and the crystallisation temperature of not more than 50°C, the process comprising homopolymerising propylene in the presence of a metallocene catalyst of general formula:



wherein C_p is a substituted cyclopentadienyl ring; C_p' is a substituted or unsubstituted fluorenyl ring; R'' is a structural bridge imparting stereorrigidity to the component; R_1 is a substituent on the cyclopentadienyl ring which is distal to the bridge, which distal substituent comprises a bulky group of the formula XR_a^* , in which X is chosen from Group IVA, $a=2$, and one R^* is chosen from hydrogen or hydrocarbyl of from 1 to 20 carbon atoms and the other different R^* is chosen from a substituted or unsubstituted cycloalkyl where X is a carbon atom in the cycloalkyl ring, R_2 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and positioned *non-vicinal* to the distal substituent and is of the formula $YR_\#$, in which Y is chosen from Group IVA, and each $R_\#$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, R_3 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and is a hydrogen atom or is of the formula $ZR_\$$, in which Z is chosen from Group IVA, and each $R_\$$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, each R_n' is the same or different and is hydrocarbyl having 1 to 20 carbon atoms in which $0 \leq n \leq 8$; M is a Group IVB transition metal or vanadium and each Q is hydrocarbyl having 1 to 20 carbon atoms or is a halogen.

The metallocene catalyst may be used either alone or in a mixture of such metallocene catalysts.

In the bulky distal substituent group R_1 , X is preferably C or Si. When a is 3, R^* may be a hydrocarbyl such as alkyl, aryl, alkenyl, alkyl aryl or aryl alkyl, preferably methyl, ethyl, propyl, isopropyl, butyl, isobutyl, amyl, isoamyl, hexyl, heptyl, octyl, nonyl, decyl, cetyl or phenyl. R_1 may comprise a hydrocarbyl which is attached to a single carbon atom in the cyclopentadienyl ring or may be bonded to two carbon atoms in that ring. Preferably, R_1 is $C(CH_3)_3$. When a is 2, one R^* is a substituted or unsubstituted cycloalkyl group, with X being C and incorporated in the cycloalkyl ring. Thus R_1 may comprise an alkyl-cycloalkyl group, typically methyl-cyclohexyl.

The proximal substituents R_2 and R_3 are the same or different and preferably CH_3 or hydrogen.

R'' is preferably isopropylidene in which the two C_p rings are bridged at position 2 of the isopropylidene.

M is preferably zirconium.

Q is preferably a halogen and more preferably Cl.

The fluorenyl ring C_p' can have up to 8 substituent groups R'_n , each of which is the same or different and may be hydrogen or a hydrocarbyl selected from alkyl, aryl, alkenyl, alkyl aryl or aryl alkyl, such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, amyl, isoamyl, hexyl, heptyl, octyl, nonyl, decyl, cetyl or phenyl. These substituents must be selected so that they do not interfere with coordination of the monomer to the metal. Preferably, therefore, the fluorenyl ring is

unsubstituted at both positions 4 and 5, these positions being distal to the bridge.

The metallocene catalyst component can be used to produce isotactic polypropylene homopolymer, optionally with a small amount of comonomer incorporated therein, i.e. a homopolymer with a low degree of copolymer. Both types of polypropylene are characterised by a low melting temperature and a high crystallisation temperature.

In accordance with a preferred aspect of the invention, the catalyst, which is selected from the cyclopentadienyl fluorenyl family, is used in a homogeneous or heterogeneous (i.e. supported catalyst) polymerisation for producing isotactic polypropylene homopolymer. In one preferred embodiment, the catalyst is a methyl cyclohexyl disubstituted cyclopentadienyl fluorenyl. Another preferred catalyst is tertiary butyl disubstituted cyclopentadienyl fluorenyl. In a further embodiment, the catalyst may be a tertiary butyl monosubstituted cyclopentadienyl fluorenyl. Such cyclopentadienyl fluorenyl metallocene catalysts enable relatively low melting points to be achieved, typically lower than 145°C for isotactic polypropylene.

By carefully selecting the catalyst formulation, isotactic polypropylene having differing melting points can be obtained without using any comonomer. A melting point as low as 120°C is achievable using these catalysts in isotactic polypropylene homopolymer. If a small concentration of comonomer for example ethylene or butene, is added to the propylene during polymerisation, a melting point lower than 120°C may be achieved.

Compared to known polypropylene copolymers or homopolymers fabricated using Ziegler-Natta catalysts, for a given melting temperature higher crystallisation temperatures are achieved

in accordance with the invention. This provides for example a significant reduction in the cycle time for injection moulding and injection and extrusion blow moulding and higher line speeds for production of film, together with reduced stickiness of the films and higher production speeds for pipe, tube and profile extrusion. The selection of a cyclopentadienyl metallocene catalyst also provides improved mechanical properties, in particular flexural and tensile properties, for the isotactic polypropylene. Furthermore, the polypropylenes produced in accordance with the invention have good light transmittance, including high transparency and low haze.

When the polymers produced in accordance with the invention are employed to produce articles by a processing technique, such as injection moulding, injection or extrusion blow moulding, or production of films, or extrusion of pipes, tubes or profiles, the polypropylene can be employed either in its pure form, or in a blend. When used pure, the polypropylene can be a layer of a multilayer or any other type of construction.

In accordance with the invention, the provision of an isotactic polypropylene homopolymer, optionally with a small degree of copolymer therein, provides not only a relatively low melting temperature of the polymer, but also a high crystallisation temperature. This in turn reduces the temperature "window" between those two temperatures to typically less than about 50°C. This greatly improves the processability of the polypropylene polymer since not only is it possible to process the polymer melt at lower temperatures, but also the speed of solidification of the polypropylene from the melt is enhanced as a result of the increased crystallisation temperature for a given melt temperature.

When the catalyst is a tertiary butyl monosubstituted cyclopentadienyl fluorenyl catalyst, the catalyst may comprise

isopropylidene (3-tert butyl-cyclopentadienyl-fluorenyl) ZrCl_2 . The production of such a catalyst is disclosed in EP-A-0537130. This catalyst produces isotactic polypropylene of low isotacticity, typically having 75 to 80% mmmm and a melting point of around 127 to 129°C. The polymer produced has a low molecular weight typically ranging from about 50,000 to 75,000 Mw.

When the catalyst comprises a tertiary butyl disubstituted cyclopentadienyl fluorenyl metallocene catalyst, the catalyst may particularly comprise isopropylidene (3-tert butyl-5-methyl cyclopentadienyl-fluorenyl) ZrCl_2 . The synthesis of such a catalyst is disclosed in EP-A-0881236. This catalyst has a higher productivity than the corresponding monosubstituted catalyst described above, and the polymer produced has a higher molecular weight, a higher isotacticity and a higher melting point. Typically, the tacticity ranges from about 83 to 86% mmmm and the melting point ranges from about 139 to 144°C. With the additional hydrocarbyl group at the five position in the cyclopentadienyl group, the amount of regio defects is reduced below the limit of detection of the NMR, i.e. to less than 0.1% regio defects.

The preferred methyl cyclohexyl disubstituted catalyst incorporates a cycloalkyl group at the three-position of the cyclopentadienyl ring. The synthesis of the methyl cyclohexyl disubstituted cyclopentadienyl fluorenyl is similar to that of the tertiary butyl disubstituted cyclopentadienyl fluorenyl catalyst but cyclohexane is used instead of acetone in the synthesis of the fulvene. Like the tertiary butyl disubstituted catalyst, the methyl cyclohexyl disubstituted catalyst produces isotactic polypropylene homopolymers having a high degree of tacticity, where mmmm is typically around 80% and a low level of regio defects, typically less than 0.1%. The melting point is around 140°C. Generally, the isotacticity and the melting point are slightly lower for the methyl

cyclohexyl disubstituted catalyst as compared to the tertiary butyl disubstituted catalyst.

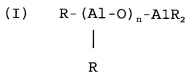
The catalyst may be a trisubstituted cyclopentadienyl fluorenyl catalyst, with for example tertiary butyl or an alkyl-cycloalkyl group at the three-position of the cyclopentadienyl ring.

When copolymerisation is selected for lower the melting temperature, metallocene catalysts offer two significant advantages over the Ziegler-Natta catalysts. First, the insertion of comonomer is more periodic, for all chain lengths, and therefore less comonomer is required for a given decrease in the melting temperature. Less comonomer is thus necessary in order to reach the desired low melting temperature. The use of comonomer to lower the melting point is thus more efficient. Second, the melting temperature of the isotactic homopolymers of propylene produced with metallocene catalysts is lower than that of the homopolymers obtained with the Ziegler-Natta catalysts. The metallocene catalysts used in accordance with the invention are Cp-Fluorenyl metallocene catalysts which produce homopolymers with melting temperatures of below 145°C.

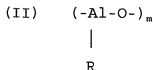
The catalyst system for use in preparing isotactic polypropylene comprises (a) a catalyst component as defined above; and (b) an aluminium- or boron-containing cocatalyst capable of activating the catalyst component. Suitable aluminium-containing cocatalysts comprise an alumoxane, an alkyl aluminium and/or a Lewis acid.

The aluminoxanes usable as such cocatalysts are well known and preferably comprise oligomeric linear and/or cyclic alkyl alumoxanes represented by the formula:

11



for oligomeric, linear alumoxanes and



for oligomeric, cyclic alumoxanes, wherein n is 1-40, preferably 10-20, m is 3-40, preferably 3-20 and R is a C₁-C₈ alkyl group and preferably methyl. Generally, in the preparation of alumoxanes from, for example, aluminium trimethyl and water, a mixture of linear and cyclic compounds is obtained.

Suitable boron-containing cocatalysts may comprise a triphenylcarbenium boronate such as tetrakis-pentafluorophenyl-borato-triphenylcarbenium as described in EP-A-0427696, or those of the general formula [L⁺-H] + [B Ar₁ Ar₂ X₃ X₄]- as described in EP-A-0277004 (page 6, line 30 to page 7, line 7).

The catalyst system may be employed in a solution polymerisation process, which is homogeneous, or a slurry process which is heterogeneous. In a solution process, typical solvents include hydrocarbons with 4 to 7 carbon atoms such as heptane, toluene or cyclohexane. In a slurry process it is necessary to immobilise the catalyst system on an inert support, particularly a porous solid support such as talc, inorganic oxides and resinous support materials such as polyolefin. Preferably, the support material is an inorganic oxide in its finely divided form.

Suitable inorganic oxide materials which are desirably employed include Group 2a, 3a, 4a or 4b metal oxides such as

silica, alumina and mixtures thereof. Other inorganic oxides that may be employed either alone or in combination with the silica, or alumina are magnesia, titania, zirconia, and the like. Other suitable support materials, however, can be employed for example, finely divided functionalised polyolefins such as finely divided polyethylene.

Preferably, the support is a silica having a surface area comprised between 200 and 700 m²/g and a pore volume comprised between 0.5 and 3 ml/g.

The amount of alumoxane and metallocenes usefully employed in the preparation of the solid support catalyst can vary over a wide range. Preferably the aluminium to transition metal mole ratio is in the range between 1:1 and 100:1, preferably in the range 5:1 to 50:1.

The order of addition of the metallocenes and alumoxane to the support material can vary. Alumoxane dissolved in a suitable inert hydrocarbon solvent may be added to the support material slurried in the same or other suitable hydrocarbon liquid and thereafter a mixture of the metallocene catalyst component is added to the slurry.

Preferred solvents include mineral oils and the various hydrocarbons which are liquid at reaction temperature and which do not react with the individual ingredients. Illustrative examples of the useful solvents include the alkanes such as pentane, iso-pentane, hexane, heptane, octane and nonane; cycloalkanes such as cyclopentane and cyclohexane, and aromatics such as benzene, toluene, ethylbenzene and diethylbenzene.

Preferably the support material is slurried in toluene and the metallocene and alumoxane are dissolved in toluene prior to addition to the support material.

The present invention will now be described in greater detail with reference to the following non-limiting Example.

Example 1

An isotactic polypropylene homopolymer was produced using, as a metallocene catalyst, isopropylidene (3-methylcyclohexyl-5-methyl cyclopentadienyl-fluorenyl)ZrCl₂.

The polymerisation was performed in a bench liquid full loop reactor in the slurry phase by introducing the metallocene catalyst precontacted with MAO (methylaluminoxane). The catalyst was unsupported. The polymerisation temperature was 60°C. The productivity of the catalyst was 65,000gPP/gcat/h. The fluff was stabilised with conventional antioxidants and then extruded and pelletised before being injection moulded into bars.

Isotactic polypropylene having a melt temperature of 142°C was obtained. The polypropylene obtained was monomodal. The isotactic polypropylene had a tacticity of 80% mmmm, with less than 0.1% regio defects. The tacticity was determined by NMR analysis and the remaining NMR results are shown in Table 1.

The polymer was also subjected to differential scanning calorimetry (DSC) to determine the melting temperature T_m and the crystallisation temperature T_c of the polypropylene. The results are shown in Table 2.

The polypropylene was also tested to determine its melt index MI₂. The melt index was determined using the procedure of ASTM-A-1238 using a load of 2.16kg at a temperature of 190°C. The melt index MI₂ of the polypropylene was 1.5g/10min.

In addition, the flexural modulus of the polypropylene was determined using the procedures of ISO R178 and the results are shown in Table 2.

Comparative Examples 1 and 2

As a comparison to the polypropylene homopolymer produced in accordance with the invention, the corresponding properties indicated in Table 2 for Example 1 were determined for a known random PP copolymer produced using a Ziegler-Natta catalyst with roughly the same melting point and MFI as for the polypropylene of Example 1 (Comparative Example 1) and for a known PP homopolymer using a Ziegler-Natta catalyst (Comparative Example 2). The sample of Comparative Example 1 was a random copolymer containing about 3.5wt% of ethylene monomer.

A comparison of the results for Example 1 and Comparative Examples 1 and 2 shows that the melting point of the isotactic polypropylene homopolymer is slightly larger than that of the random polypropylene containing about 3.5wt% of ethylene. However, in accordance with the invention the crystallisation temperature of the isotactic polypropylene is much higher, about 12°C higher, than that of the random polypropylene. The isotactic polypropylene has a difference between T_m and T_c of less than 50°C. This is a significant reduction in the temperature difference that can result in a dramatic improvement in processing performance, leading to a decrease of cycle time for injection moulding and injection and extrusion blow moulding, an increase in the line speed and a reduction in the stickiness of the film for extruding and blowing films, and an increase in the production speed for pipe, tube and profile extrusion.

For the polypropylene homopolymer of Comparative Example 2, the melting temperature is around 163°C which is higher than for Example 1 and the crystallisation temperature is also higher than for Example 1 and Comparative Example 1, being around 100°C. This yields a temperature window between the melting temperature and the crystallisation temperature which

is more than 60°C, i.e.. significantly broader than that achievable in accordance with the invention.

The flexural modulus of the isotactic polypropylene is about 20% larger than that of the random polypropylene of Comparative Example 1. For the polypropylene homopolymer of Comparative Example 2, this had a higher flexural modulus as compared to the polymer of Example 1. However the homopolymer of Comparative Example 2 has lower transparency and requires higher processing temperatures than the isotactic polymer of Example 1.

TABLE 1

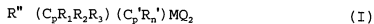
mmmm	80.38
mmmr	6.04
rmmr	1.91
mmrr	5.43
rmrr+mrmm	0.88
mrmr	0.33
rrrr	1.30
mr rr	1.13
mr rm	2.61
mm	88.32
mr	6.64
rr	5.04
r	8.36

TABLE 2

	Units	Example 1	Comparative Example 1	Comparative Example 2
MI2	g/10min	1.5	3.9	13.4
Tm	°C	142.1	138.7	163.1
Tc	°C	96.3	84.3	100.6
FLEXURAL MODULUS				
Flex. Mod. at 0.25%	MPa	904	749	1209
Flex. Mod. at 1%	MPa	868	718	1172

CLAIMS:

1. Use of isotactic polypropylene homopolymers or copolymers in processes in which the polypropylene solidifies from a melt, wherein for enhanced speed of solidification of the polypropylene the polypropylene has a melt temperature and a crystallisation temperature not more than 50°C less than the melt temperature resulting from the polypropylene having been produced using a metallocene catalyst component having the general formula:



wherein C_p is a substituted cyclopentadienyl ring; C_p' is a substituted or unsubstituted fluorenyl ring; R'' is a structural bridge imparting stereorigidity to the component; R_1 is a substituent on the cyclopentadienyl ring which is distal to the bridge, which distal substituent comprises a bulky group of the formula XR^*_a in which X is chosen from Group IVA, and when $a=3$ each R^* is the same or different and chosen from hydrogen or hydrocarbyl of from 1 to 20 carbon atoms, or when $a=2$ one R^* is chosen from hydrogen or hydrocarbyl of from 1 to 20 carbon atoms and the other different R^* is chosen from a substituted or unsubstituted cycloalkyl where X is a carbon atom in the cycloalkyl ring, R_2 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and positioned *non-vicinal* to the distal substituent and is hydrogen or of the formula $YR^\#_3$ in which Y is chosen from Group IVA, and each $R^\#$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, R_3 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and is a hydrogen atom or is of the formula $ZR^\$_3$ in which Z is chosen from Group IVA, and each $R^\$$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, each R'_n is the same or different and is hydrocarbyl having 1 to 20 carbon atoms in which $0 \leq n \leq 8$; M is a

Group IVB transition metal or vanadium and each Q is hydrocarbyl having 1 to 20 carbon atoms or is a halogen.

2. Use according to claim 1, wherein R_1 is a methylcyclohexyl group.

3. Use according to claim 1 wherein R_1 is a tertiary butyl group.

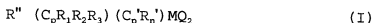
4. Use according to claim 2 or claim 3 wherein R_2 is a methyl group.

5. Use according to claim 2 or claim 3 wherein R_2 is hydrogen.

6. Use according to any foregoing claim wherein each R' is hydrogen.

7. Use according to any foregoing claim wherein Y is carbon.

8. A process for producing an isotactic homopolymer of propylene having a melt temperature of from 139 to 144°C and a difference between the melt temperature and the crystallisation temperature of not more than 50°C, the process comprising homopolymerising propylene in the presence of a metallocene catalyst of general formula:



wherein C_p is a substituted cyclopentadienyl ring; C_p' is a substituted or unsubstituted fluorenyl ring; R'' is a structural bridge imparting stereorrigidity to the component; R_1 is a substituent on the cyclopentadienyl ring which is distal to the bridge, which distal substituent comprises a bulky group of the formula XR^*_a in which X is chosen from Group IVA, $a=2$, and one R^* is chosen from hydrogen or hydrocarbyl of from 1 to

20 carbon atoms and the other different R^* is chosen from a substituted or unsubstituted cycloalkyl where X is a carbon atom in the cycloalkyl ring, R_2 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and positioned *non-vicinal* to the distal substituent and is of the formula $YR_{\#}$, in which Y is chosen from Group IVA, and each $R_{\#}$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, R_3 is a substituent on the cyclopentadienyl ring which is proximal to the bridge and is a hydrogen atom or is of the formula $ZR_{\$}$, in which Z is chosen from Group IVA, and each $R_{\$}$ is the same or different and chosen from hydrogen or hydrocarbyl of 1 to 7 carbon atoms, each R'_n is the same or different and is hydrocarbyl having 1 to 20 carbon atoms in which $0 \leq n \leq 8$; M is a Group IVB transition metal or vanadium and each Q is hydrocarbyl having 1 to 20 carbon atoms or is a halogen.

9. A process according to claim 8 wherein R_1 is a methylcyclohexyl group.

10. A process according to claim 9 wherein R_2 is a methyl group.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/EP 00/01736

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C08F10/06 C08F4/64

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 742 227 A (FINA TECHNOLOGY) 13 November 1996 (1996-11-13) cited in the application	1,6,8
Y	examples 1,2	1,3,4,6, 7
Y	page 5, line 57 -page 6, line 2	1,2,5,6
Y	examples 1,2	1-3,5,6, 8,9
	--- -/-	

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"B" document member of the same patent family

Date of the actual completion of the international search

15 June 2000

Date of mailing of the international search report

06/07/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Gamb, V

INTERNATIONAL SEARCH REPORT

Int. onal Application No

PCT/EP 00/01736

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	OBORA, YASUSHI ET AL: "Ancillary Ligand Effects in Chiral C1-Symmetric ansa-Metallocene Catalyst for Stereoregular.alpha.-Olefin Polymerization. "Wingspan" Modification with Octahydrofluorene" ORGANOMETALLICS (1997), 16(12), 2503-2505, XP002140298	8,9
Y	the whole document	1,2,5,6
X	EP 0 881 236 A (FINA RESEARCH) 2 December 1998 (1998-12-02) cited in the application page 7; table 1	8
Y	examples 1,2,4	1,3,4,6,7
Y	WO 98 06760 A (DOW CHEMICAL COMPANY, USA;NORTHWESTERN UNIVERSITY) 19 February 1998 (1998-02-19) page 4, line 24 -page 5, line 24 example 1 page 21; table 1	1-3,5,6,8,9
X	EP 0 870 779 A (FINA TECHNOLOGY) 14 October 1998 (1998-10-14) cited in the application page 5, line 2 - line 20 page 7, line 5 - line 10 page 11; example 14; table IV	1
P,X	EP 0 905 173 A (FINA TECHNOLOGY) 31 March 1999 (1999-03-31) page 2, line 5 - line 6 page 7, line 29 page 8; table 1	1
A	EP 0 537 130 A (FINA TECHNOLOGY) 14 April 1993 (1993-04-14) cited in the application examples 1-4 page 9; table 1	8

INTERNATIONAL SEARCH REPORT

Information on patent family members

Inter. Appl. Application No

PCT/EP 00/01736

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0742227 A	13-11-1996	US 5036034 A AT 189226 T CA 2027124 A CN 1051735 A, B CN 1100733 A DE 69033435 D EP 0423101 A ES 2142303 T JP 3193796 A KR 181495 B	30-07-1991 15-02-2000 11-04-1991 29-05-1991 29-03-1995 02-03-2000 17-04-1991 16-04-2000 23-08-1991 15-05-1999
EP 0881236 A	02-12-1998	WO 9854230 A EP 0984989 A	03-12-1998 15-03-2000
WO 9806760 A	19-02-1998	AU 3904297 A CA 2259428 A CN 1226255 A CZ 9900381 A EP 0917543 A	06-03-1998 19-02-1998 18-08-1999 12-05-1999 26-05-1999
EP 0870779 A	14-10-1998	JP 10338703 A	22-12-1998
EP 0905173 A	31-03-1999	JP 11166058 A	22-06-1999
EP 0537130 A	14-04-1993	US 5416228 A CA 2077713 A DE 69213878 D DE 69213878 T JP 6122718 A	16-05-1995 08-04-1993 24-10-1996 13-02-1997 06-05-1994